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**Amendments to the Claims** 

This listing of claims will replace all prior versions, and listings, of the claims in the

application.

**Claims Listing** 

1. (Currently Amended) A method comprising:

placing a sample in contact with a device, wherein at least a portion of the

device comprises a plurality of zones, wherein at least one zone is a detection zone capable

of being separated from said plurality of zones and the remainder of the device; wherein the

detection zone comprises an immobilized binding partner for an analyte and wherein binding

between the immobilized binding partner and a suspected analyte causes formation of a

detectable signal and detection of the signal indicates the presence of a suspected analyte in

the sample;

separating at least part of the portion of the device detection zone containing

the bound analyte and immobilized binding partner from said plurality of zones and the

remainder of the device; and

analyzing the portion containing the bound analyte and immobilized binding

partner without detaching the bound analyte from the immobilized binding partner to provide

information regarding the suspected analyte.

2. (Previously Presented) The method of Claim 1, wherein the device is a lateral

flow device.

3. (Previously Presented) The method of Claim 1, wherein the information

identifies the suspected analyte.

4. (Previously Presented) The method of Claim 1, wherein the information

describes one or more characteristics of the suspected analyte.

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5. (Previously Presented) The method of Claim 1, further comprising:

placing the portion containing the bound analyte and immobilized binding partner in conditions effective to cause the quantity of the suspected analyte to increase; and separating at least part of the suspected analyte from the portion containing the bound analyte and immobilized binding partner after an increase in the quantity of the

suspected analyte.

6. (Previously Presented) The method of Claim 1, wherein the analyte is an organism and analyzing the portion containing the bound analyte and immobilized binding partner comprises placing the portion containing the bound analyte and immobilized binding partner on or in a selective growth medium in which the analyte will proliferate if present.

7. (Currently Amended) The method of Claim 1, wherein the method further comprises storing the device without further processing for up to five days after placing the sample in contact with the device and before separating the portion of the device detection zone containing the bound analyte and immobilized binding partner from the plurality of zones and the remainder of the device.

8. (Withdrawn) A method for isolating or concentrating a substance consisting essentially of:

placing a substance in contact with a device, wherein at least a portion of the device comprises an immobilized binding partner for an analyte and wherein binding between the immobilized binding partner and a suspected analyte causes formation of a detectable signal and detection of the signal indicates the presence of the suspected analyte in the substance;

separating the portion of the device containing the bound analyte and immobilized binding partner from the remainder of the device; and

analyzing the portion containing the bound analyte and immobilized binding partner to provide information regarding the suspected analyte.

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9. (Currently Amended) A kit for performing the method of claim 1, comprising

a device; wherein at least a portion of the device comprises a detection zone plurality of

zones, wherein at least one zone is a detection zone; wherein the detection zone comprises an

immobilized binding partner for an analyte, and is capable of being separated from said

plurality of zones and the remainder of the device and analyzed to provide information

regarding the bound analyte.; and wherein the detection zone or a portion thereof is separable

from the remainder of the device.

10. (Currently Amended) A device wherein at least a portion of the device

comprises a plurality of zones, wherein at least one zone is a detection zone capable of being

separated from said plurality of zones and the remainder of the device; detection zone,

wherein the detection zone comprises an immobilized binding partner for an analyte; and

wherein binding between the immobilized binding partner and a suspected analyte causes

formation of a detectable signal in the detection zone, and wherein the device comprises

structural features that facilitate separation of the detection zone containing the bound

analyte and the immobilized binding partner or a part of the detection zone containing the

bound analyte and the immobilized binding partner from the plurality of zones and the

remainder of the device., wherein said detection zone or part thereof can be analyzed to

provide information regarding the bound analyte.

11. (Withdrawn) The method of claim 8, wherein the substance is a food or soil

contaminant.

12. (Withdrawn) The method of claim 8, wherein the substance is a

microorganism.

13. (Withdrawn) The method of claim 8, wherein the substance is a pathogen.

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14. (Withdrawn) The method of Claim 8, wherein the device is a lateral flow

device.

15. (Withdrawn) The method of Claim 8, wherein the information identifies the

suspected analyte.

16 (Withdrawn) The method of Claim 8, wherein the information describes one or

more characteristics of the suspected analyte.

17. (Withdrawn) The method of Claim 8, further comprising:

placing the portion containing the bound analyte and immobilized binding

partner in conditions effective to cause the quantity of the suspected analyte to increase; and

separating at least part of the suspected analyte from the portion containing the

bound analyte and immobilized binding partner after an increase in the quantity of the

suspected analyte.

18. (Withdrawn) The method of Claim 8, wherein the analyte is an organism and

analyzing the portion containing the bound analyte and immobilized binding partner

comprises placing the portion containing the bound analyte and immobilized binding partner

on or in a selective growth medium in which the analyte will proliferate if present.

19. (Withdrawn) The method of Claim 8, wherein the method further comprises

storing the device without further processing for up to five days after placing the sample in

contact with the device and before separating the portion of the device containing the bound

analyte and immobilized binding partner from the reminder of the device.

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20. (Previously Presented) The method of Claim 10, wherein the device is a lateral

flow device.

21. (Withdrawn) The method of claim 8, wherein analyzing the portion containing

the bound analyte and immobilized binding partner to provide information regarding the

suspected analyte comprises analyzing the portion containing the bound analyte and

immobilized binding partner using a strip test binding assay, an agglutination assay, a DNA

polymerase chain reaction test, a motility test, a toxicology test, serotyping, selective media

or selective agar plating.

22. (Withdrawn) The method of claim 8, wherein analyzing the portion containing

the bound analyte and immobilized binding partner to provide information regarding the

suspected analyte comprises analyzing the portion containing the bound analyte and

immobilized binding partner using a DNA polymerase chain reaction test.

23. (Withdrawn) The method of claim 8, wherein analyzing the portion containing

the bound analyte and immobilized binding partner to provide information regarding the

suspected analyte comprises analyzing the portion containing the bound analyte and

immobilized binding partner using selective media or selective agar plating.

24. (Withdrawn) The method of claim 8, wherein the substance is Escherichia

coli, Salmonella or Listeria.

25. (Withdrawn) The method of claim 8, wherein the substance is Escherichia coli

O157:H7.